

PCT**INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY**

(Chapter II of the Patent Cooperation Treaty)

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference A03P2012P	FOR FURTHER ACTION		See Form PCT/PEA/416
International application No. PCT/SE2003/001494	International filing date (day/month/year) 25.09.2003	Priority date (day/month/year) 25.09.2003	
International Patent Classification (IPC) or national classification and IPC A61N1/37			
Applicant ST. JUDE MEDICAL AB et al.			

<p>1. This report is the international preliminary examination report, established by this International Preliminary Examining Authority under Article 35 and transmitted to the applicant according to Article 36.</p> <p>2. This REPORT consists of a total of 5 sheets, including this cover sheet.</p> <p>3. This report is also accompanied by ANNEXES, comprising:</p> <ul style="list-style-type: none"> a. <input type="checkbox"/> <i>(sent to the applicant and to the International Bureau) a total of . sheets, as follows:</i> <ul style="list-style-type: none"> <input type="checkbox"/> sheets of the description, claims and/or drawings which have been amended and are the basis of this report and/or sheets containing rectifications authorized by this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions). <input type="checkbox"/> sheets which supersede earlier sheets, but which this Authority considers contain an amendment that goes beyond the disclosure in the international application as filed, as indicated in item 4 of Box No. I and the Supplemental Box. b. <input type="checkbox"/> <i>(sent to the International Bureau only) a total of (indicate type and number of electronic carrier(s)) , containing a sequence listing and/or tables related thereto, in computer readable form only, as indicated in the Supplemental Box Relating to Sequence Listing (see Section 802 of the Administrative Instructions).</i>
<p>4. This report contains indications relating to the following items:</p> <ul style="list-style-type: none"> <input checked="" type="checkbox"/> Box No. I Basis of the opinion <input type="checkbox"/> Box No. II Priority <input type="checkbox"/> Box No. III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability <input type="checkbox"/> Box No. IV Lack of unity of invention <input checked="" type="checkbox"/> Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement <input type="checkbox"/> Box No. VI Certain documents cited <input checked="" type="checkbox"/> Box No. VII Certain defects in the international application <input type="checkbox"/> Box No. VIII Certain observations on the international application

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Name and mailing address of the international preliminary examining authority:  European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d Fax: +49 89 2399 - 4465	Authorized Officer Kempin, H-F Telephone No. +49 89 2399-2716



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Box No. I Basis of the report

1. With regard to the **language**, this report is based on the international application in the language in which it was filed, unless otherwise indicated under this item.
 - This report is based on translations from the original language into the following language, which is the language of a translation furnished for the purposes of:
 - international search (under Rules 12.3 and 23.1(b))
 - publication of the international application (under Rule 12.4)
 - international preliminary examination (under Rules 55.2 and/or 55.3)
2. With regard to the **elements*** of the international application, this report is based on (*replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report*):

Description, Pages

1-5 as published

Claims, Numbers

1-4 as published

Drawings, Sheets

1/5-5/5 as published

a sequence listing and/or any related table(s) - see Supplemental Box Relating to Sequence Listing

3. The amendments have resulted in the cancellation of:
 - the description, pages
 - the claims, Nos.
 - the drawings, sheets/figs
 - the sequence listing (*specify*):
 - any table(s) related to sequence listing (*specify*):
4. This report has been established as if (some of) the amendments annexed to this report and listed below had not been made, since they have been considered to go beyond the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)).
 - the description, pages
 - the claims, Nos.
 - the drawings, sheets/figs
 - the sequence listing (*specify*):
 - any table(s) related to sequence listing (*specify*):

* *If item 4 applies, some or all of these sheets may be marked "superseded."*

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Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)	Yes:	Claims	1-4
	No:	Claims	
Inventive step (IS)	Yes:	Claims	1-4
	No:	Claims	
Industrial applicability (IA)	Yes:	Claims	1-4
	No:	Claims	

2. Citations and explanations-(Rule 70.7):

see separate sheet

Box No. VII Certain defects in the international application

The following defects in the form or contents of the international application have been noted:

see separate sheet

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Re Item V

Reasoned statement with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

Reference is made to the following documents:

D1: EP-A-1 155 712

The document D1 is regarded as being the closest prior art to the subject-matter of claim 1, and shows (the references in parentheses applying to this document):

A biventricular stimulation device (see 10, 32, 48 in connection with column 8, lines 22-27) comprising a pulse generator for delivering stimulation pulses at least to the ventricles of the heart (see 72a, 72 b in figure 1), and an evoked response detector having independent first and second ventricular sensing channels for ventricular evoked response detection of the ventricles (see 84a, 84b and column 17, lines 38, 39), said pulse generator being controlled to deliver a stimulation pulse to the second ventricle with a VV time delay after stimulation pulse delivery to the first stimulated ventricle (see column 23, lines 40-43) which is shorter than an evoked response detection time window following delivery of said stimulation pulse to the first stimulated ventricle (see column 13, lines 16-21 and column 23, line 53 to column 24, line 4).

The subject-matter of claim 1 differs from this known biventricular stimulation device by the features of the second part of claim 1.

The subject-matter of claim 1 is therefore new (Article 33(2) PCT).

The problem to be solved by the present invention may be regarded as to improve detection of evoked response in a biventricular pacemaker.

The solution to this problem proposed in claim 1 of the present application is considered as involving an inventive step (Article 33(3) PCT) for the following reasons:

Document D1 detects the evoked response by using the far field signal, e.g. when the right ventricle is paced the evoked response is detected in the left ventricular sense channel. Hence, the second ventricle cannot be paced when the evoked response detection time

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window is open after pacing of the first ventricle. The other documents cited in the International Search Report do not incite the skilled person to modify the pacer known from D1 as defined in the second part of claim 1 since they do not disclose any details of evoked response detection in connection with biventricular pacing with short VV delay.

Claims 2-4 are dependent on claim 1 and as such also meet the requirements of the PCT with respect to novelty and inventive step.

Re Item VII

1. The features of the claims are not provided with reference signs placed in parentheses (Rule 6.2(b) PCT).
2. A document reflecting the prior art described on page 1, first paragraph (e.g. document D1) is not identified in the description (Rule 5.1(a)(ii) PCT).